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Jc514 U.S. PTO

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PROVISIONAL APPLICATION FOR PATENT COVER SHEET
(Large Entity)

A/prov

This is a request for filing a PROVISIONAL APPLICATION FOR PATENT under 37 CFR 1.53 (b)(2).

Docket Number		1038.766 MIS		Type a plus sign (+) inside this box	+
INVENTOR(S)/APPLICANT(S)					
LAST NAME	FIRST NAME	MIDDLE INITIAL	RESIDENCE (CITY AND EITHER STATE OR FOREIGN COUNTRY)		
PARRINGTON	Mark		45 Martin Street, Bradford, Ontario, Canada, L3Z 1Z4.		
TITLE OF THE INVENTION (200 characters max)					
ALPHAVIRUS VECTORS					
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STATE	Ontario	ZIP CODE	MSG 1R7	COUNTRY	Canada
ENCLOSED APPLICATION PARTS (check all that apply)					
<input checked="" type="checkbox"/>	Specification	Number of Pages	8		
<input checked="" type="checkbox"/>	Drawing(s)	Number of Sheets	7	<input type="checkbox"/> Other (specify)	
METHOD OF PAYMENT OF FILING FEES FOR THIS PROVISIONAL APPLICATION FOR PATENT (check one)					
<input checked="" type="checkbox"/>	A check or money order is enclosed to cover the filing fees			FILING FEE AMOUNT	\$150.00
<input type="checkbox"/>	The Commissioner is hereby authorized to charge filing fees and credit Deposit Account Number:				

The invention was made by an agency of the United States Government or under a contract with an agency of the United States

☒ No.☐ Yes, the name of the U.S. Government agency and the Government contract number

Respectfully submitted,

SIGNATURE Michael I. Stewart

Date 11/14/1997

TYPED or PRINTED NAME Michael I. Stewart

REGISTRATION NO. (if appropriate) 24,973

☐ Additional inventors are being named on separately numbered sheets attached hereto**USE ONLY FOR FILING A PROVISIONAL APPLICATION FOR PATENT**
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P18LARGE/REV03

MIS 1038-766 1997 11 14 D1

TITLE OF INVENTIONALPHAVIRUS VECTORS

5

FIELD OF INVENTION

The present invention relates to the field of DNA vaccines and is particularly concerned with modified alpha virus vectors for use in such vaccines.

BACKGROUND OF THE INVENTION

10 Semliki Forest virus (SFV) is a member of the Alphavirus genus in the Togaviridae family. The mature virus particle contains a single copy of a ssRNA genome with a positive polarity that is 5'-capped and 3'-polyadenylated. It functions as an mRNA and naked RNA
15 can start an infection when introduced into cells. Upon infection/transfection, the 5' two-thirds of the genome is translated into a polyprotein that is processed into the four nonstructural proteins (nsP1 to 4) by self cleavage. Once the ns proteins have been synthesized
20 they are responsible for replicating the plus-strand (42S) genome into full-length minus strands (ref. 14). These minus-strands then serve as templates for the synthesis of new plus-strand (42S) genomes and the 26S subgenomic mRNA (ref. 1 - Throughout this application,
25 various references are cited in parentheses to describe more fully the state of the art to which this invention pertains. Full bibliographic information for each citation is found at the end of the specification. The disclosures of these references are hereby incorporated
30 by reference into the present disclosure). This subgenomic mRNA, which is colinear with the last one-third of the genome, encodes the SFV structural proteins. In 1991 Liljestrom and Garoff (ref. 2) designed a series of expression vectors based on the SFV
35 cDNA replicon. These vectors had the virus structural protein genes deleted to make the way for heterologous

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inserts, but preserved the nonstructural coding region for production of the nsP1 to 4 replicase complex. Short 5' and 3' sequence elements required for RNA replication were also preserved. A polylinker site was
5 inserted downstream from the 26S promoter followed by translation stop sites in all three frames. An SpeI site was inserted just after the 3' end of the SFV CDNA for linearization of the plasmid for use in vitro transcription reactions.

10 Injection of SFV RNA encoding a heterologous protein have been shown to result in the expression of the foreign protein and the induction of antibody in a number of studies (refs. 3,4). The use of SFV RNA inoculation to express foreign proteins for the purpose
15 of immunization would have several of the advantages associated with plasmid DNA immunization. For example, SFV RNA encoding a viral antigen may be introduced in the presence of antibody to that virus without a loss in potency due to neutralization by antibodies to the
20 virus. Also, because the protein is expressed in vivo the protein should have the same conformation as the protein expressed by the virus itself. Therefore, concerns about conformational changes which could occur during protein purification leading to a loss in
25 immunogenicity, protective epitopes and possibly immunopotential, could be avoided by plasmid DNA immunization.

In WO95/27044, the disclosure of which is incorporated herein by reference, there is described the
30 use of alphavirus cDNA vectors based on cDNA complementary to the alphavirus RNA sequence. Once transcribed from the cDNA under transcriptional control of a heterologous promoter, the alphavirus RNA is able to self-replicate by means of its own replicase and
35 thereby amplify the copy number of the transcribed recombinant RNA molecules.

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SUMMARY OF THE INVENTION

5 The present invention is concerned with modifications to the alphavirus cDNA vectors described in the aforementioned WO 95/27044 to permit enhanced replication of the alphavirus. In the present invention, a heterologous splice site is introduced into the alphavirus replicon sequence, particularly that of Semliki Forest virus (SFV).

10 Accordingly, in one aspect, the present invention provides a cDNA molecule complementary to at least part of an alphavirus RNA genome, which cDNA molecule comprises the complement of the complete alphavirus RNA genome regions which are essential for replication of the said alphavirus RNA, and further comprises a
15 heterologous cDNA sequence capable of expression in an animal or human host cell, said heterologous cDNA sequence being inserted into a region of the cDNA molecule which is non-essential to replication thereof, and the cDNA molecule being placed under
20 transcriptional control of a promoter sequence functional in said animal or human cell, wherein at least one heterologous splice site is provided in the complement of the complete alphavirus RNA genome regions which are essential for replication of the
25 alphavirus RNA, to prevent aberrant RNA splicing of the alphavirus.

The alphavirus molecule is a large molecule and, accordingly, there is a high probability of splice sites, thereby impairing the replication of the
30 alphavirus and hence its ability to express the heterologous DNA is impaired. By introducing the at least one heterologous splice site in accordance with the present invention into the alphavirus replicon sequence, any splicing is likely to be directed at the
35 heterologous splice site rather than any endogenous splice site.

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In the constructs provided herein, the promoter may be directly coupled to the 5'-end of the alphavirus sequence, which has the effect of reducing the variability in the splicing event at the 5'- end of the alphavirus replicon.

In addition, there may be provided at the 3'end of the Simliki Forest virus segment, a hepatitis delta ribosyme sequence to ensure proper *in vivo* cleavage at the 3'-end of the sequence. Any other convenient sequence may be employed to achieve this effect.

The heterologous splice site sequence may be provided by the nucleotide sequence of the rabbit β -globin intron II, as described in reference 5. Such heterologous splice site sequence may be inserted into the complement sequence at any convenient location which does not preclude replication of the alphavirus.

I have identified five suitable sites in the SFV replicon, which are contained within an EcoRV-SpeI fragment of the replicon which is 7983 bp in length (Fig. 3). The first such site is a Ppu-MI site, at position 2719 within the EcoRV-SpeI fragment.

In constructing the modified vectors provided herein, the EcoRV-SpeI fragment is cut with Ppu-MI at position 2719 and made blunt-ended with Mung Bean nuclease, which removes three bases from the SFV sequence. A blunt-ended β -globin II intron, which is 536 bp long, is ligated into the site and replaces the missing three bases with sequence added to the 3'-end of the β -globin intron sequence (Fig. 1).

The other four suitable sites for insertion of the Intron are the PvuII sites at bp 2518 3113, 6498 and 6872 of the EcoRV-SpeI fragment. Insertion of the Intron is achieved by cutting with PvuII (a blunt end cutter) and the blunt-ended β -globin II intron sequence (Fig. 2) is ligated into one or more of these sites.

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BRIEF DESCRIPTION OF DRAWINGS

Figure 1 shows the DNA sequence of the β -globin II intron encoding three additional nucleotides at the 3'-end thereof (SEQ ID No:1);

Figure 2 shows the DNA sequence of the β -globin II intron (SEQ ID No:2); and

Figures 3A to 3E show the DNA sequence of the EcoRV-SpeI fragment of Semliki Forest virus replicon (SEQ ID No:3).

GENERAL DESCRIPTION OF INVENTION

As discussed above, the present invention provides a modified alphavirus cDNA. The alphavirus preferably is Semliki Forest virus.

The promoter sequence may comprise a promoter of eukaryotic or prokaryotic origin. Suitable promoters are the cytomegalovirus immediate early promoter (pCMV), although other promoters, such as the Rous sarcoma virus long-terminal repeat promoter (pRSV), since, in the case of these and similar promoters, transcription is performed by the DNA-dependent RNA polymerase of the host cell. Additionally, the SP6, T3 or T7 promoters can be used, provided that the cell has first been transformed with genes encoding SP6, T3 or T7 RNA polymerase molecules which are either inserted into the chromosome or remain episomal. Expression of these (SP6, T3, T7) RNA polymerase-encoding genes is dependent on the host cell DNA-dependent RNA polymerase.

The heterologous cDNA insert may comprise the coding sequence for a desired product, which may be a biologically active protein or polypeptide, e.g., an immunogenic or antigenic protein or polypeptide, or a therapeutically active protein or polypeptide. The heterologous cDNA may also comprise additional

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sequences, such as a sequence complementary to an RNA sequence which is a self-cleaving ribozyme sequence.

The DNA vectors provided herein may be administered to a host, including a human host, for in vivo expression of the heterologous cDNA sequence, in accordance with a further aspect of the invention, in order to generate an immune response in the host, which may be a protective immune response. The DNA vectors may be further formulated into immunogenic compositions for such administration.

SUMMARY OF DISCLOSURE

In summary of this disclosure, the present invention provides a modified alphavirus-based expression vector wherein at least one splice site is introduced to the alphavirus replicon to prevent aberrant splicing of the alphavirus genome. Modifications are possible within the scope of the invention.

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REFERENCES

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- 5 2. Chin, J., Magoffin, R.L., Shearer, L.A., Schieble, J.H. and Lennette, E.H. (1969) Am. J. Epidemiol. 89 (4), 449-463.
- 10 3. Jensen, K.E., Peeler, B.E. and Dulworth, W.G. (1962) J. Immunol. 89, 216-226.
- 15 4. Murphy, B.R., Prince, G.A., Collins, P.L., Van Wyke-Coelingh, K., Olmstead, R.A., Spriggs, M.K., Parrott, R.H., Kim, H.-Y., Brandt, C.D. and Chanock, R.N. (1988) Vir. Res. 11, 1-15.
- 20 5. Chapman, B.S.; Thayer, R.M.; Vincent, K.A. and Haigwood, N.L., Nucl. Acids. Res. 1991, 19: 3979-3986.

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ABSTRACT OF THE DISCLOSURE

A modified alphavirus expression vector is provided wherein at least one heterologous splice site is introduced to the alphavirus replicon to prevent aborrant splicing of the alphavirus, which may be Semliki Forest virus following administration of the vector to a host.

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10 20 30 40 50 60 70 80 90 100 110 120 130 140 150 160 170 180 190 200 210 220 230 240 250 260 270 280 290 300 310 320 330 340 350 360 370 380 390 400 410 420 430 440 450 460 470 480 490 500 510 520 530 540 550 560 570 580 590 600
 GTGAGTTTGG GGACCCCTTGA TTGTTCTTTC TTTTTCGCTA TTGTAATTT CATCTTATAT
 GGAGGGGGCA AAGTTTTCAG GGTGTTGTTT AGAATGGGAA GATGTCCCTT GTATCACCAT
 GGACCCCTCAT GATAATTTTG TTTCTTTTAC TTTCTACTCT GTTGACAACC ATTGTCTCCT
 CTTATTTTCT TTTCAATTTTC TGTAACTTTT TCGTTAAACT TTAGCTTGCA TTTGTAACGA
 ATTTTAAAT TCACTTTGT TTAATTGTCA GATTGTRAGT ACTTCTCTA ATCACTTTT
 TTCAAGGCA ATCAGGGTAT ATTATATTGT ACTTCAGCAC AGTTTTAGAG AACAAATTGTT
 ATAATTAAAT GATAAGGTAG AATATTCTTG CATATAAAT CTGGCTGGCG TGGAAATATT
 CTTATTGGTA GAAACAACTA CATCTGGTC ATCATCTTGC CTTTCTCTTT ATGTTTACAA
 TGATATACAC TGTTTGAGAT GAGGATATAA TACTCTGAGT CCAACCGGG CCCCTCTGCT
 AACCATGTC ATGCCCTTCT CTTTCTCTA CAGGTC.....

B-globin +
 3 2FV bases.

B. gelatin II

Fig 2

- 1 -

10 20 30 40 50 60
GTGAGTTGG GGACCCCTGA TTGTTCTTC TTTTGGCTA TTTTAAATT CAGTTATAT
70 80 90 100 110 120
GGAGGGGCA AAGTTTTCAG GGTGTTGTTT AGAATGGGA GATGTCCCTT GTATCACCAT
130 140 150 160 170 180
GGACCCCTAT GATAAATTTC TTCTTTTCAC TTTCTACTCT GTTGACAACC ATTGTCTCCT
190 200 210 220 230 240
CTTATTTTCT TTTCAATTTTC TGTAACTTTT TCGTTAAACT TTAGCTTGCA TTTGTAAACA
250 260 270 280 290 300
ATTTTAAAT TCACCTTTTG TTATTGTCA GATTGTAAAT ACTTCTCTA ATCATTTTT
310 320 330 340 350 360
TTTCAAGGCA ATCAGGGGTAT ATTATATTGT ACTTCAGCAC AGTTTTAGAG AACAAATGTT
370 380 390 400 410 420
ATAATTAAAT GATAAGGTAG AATATTTCTG CATATAAATT CTGGCTGGCG TGGAAATATT
430 440 450 460 470 480
CTTATTGGTA GAAACAATA CATCTCTGTC ATCATCCTGC CTTTCTCTTT ATGGTTACAA
490 500 510 520 530 540
TGATATACAC TGTTTGAGAT GAGGATATAA TACTCTGAGT CCAAAACGGG CCCCTCTGCT
550 560 570 580 590 600
AACCATGTTT ATGCTTCTT CTTTTTCCTA CAG.....

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EcoRV-SpeI

Fragment

10	20	30	40	50
ATCGGCAGTG	CGCCTTCCAG	GAGATATG	TCTACGCACA	AATACCACTG
70	80	90	100	110
ATGCGCAGCG	CAGAAGACCC	CGAAAGGCTC	GATAGCTACG	CAAAGAAACT
130	140	150	160	170
TCCGGGAAGG	TGCTGGATAG	AGAGATCGCA	GGAAAAATCA	CCGACCTGCA
190	200	210	220	230
GCTACGCCAG	ACGCTGAATC	TCCTACCTTT	TGCCTGCATA	CAGACGTCAC
250	260	270	280	290
GCAGCCGAAG	TGGCCGTATA	CCAGGACGTG	TATGCTGTAC	ATGCACCAAC
310	320	330	340	350
CATCAGGCCA	TGAAAGGTGT	CAGAACGGCG	TATTGGATTG	GGTTTGACAC
370	380	390	400	410
ATGTTTGACG	CGCTAGCAGG	CGCGTATCCA	ACCTACGCCA	CAAACCTGGC
430	440	450	460	470
GTGTTACAGG	CCAGGAACAT	AGGACTGTGT	GCAGCATCCT	TGACTGAGGG
490	500	510	520	530
AAACIGTCCA	TTCTCCGCAA	GAAGCAATTG	AAACCTTGCG	ACACAGTCAT
550	560	570	580	590
GGATCTACAT	TGTACACTGA	GAGCAGAAAG	CTACTGAGGA	GCTGGCACTT
610	620	630	640	650
TTCCACCTGA	AAGGTAAACA	ATCCTTTACC	TGTAGGTGCG	ATACCATCGT
670	680	690	700	710
GGGTACGTAG	TTAAGAAAAT	CACTATGTGC	CCCGGCCTGT	ACGGTAAAC
730	740	750	760	770
GCCGTGACGT	ATCACGCGGA	GGGATTCCTA	GTGTGCAAGA	CCACAGACAC
790	800	810	820	830
GAAAGAGTCT	CATTCCCTGT	ATGCACCTAC	GTCCCCCTCA	CCATCTGTGA
850	860	870	880	890
GGGATACTAG	CGACCGACGT	CACACCGGAG	GACGCACAGA	AGTTGTTAGT
910	920	930	940	950
CAGAGGATAG	TTGTGAACGG	AAGAACACAG	CGAAACACTA	ACACGATGAA
970	980	990	1000	1010
CTTCCGATTG	TGGCCGTGCG	ATTTAGCAAG	TGGGCGAGGG	AATACAAGGC
1030	1040	1050	1060	1070
GATGAAAAAC	CTCTGGGTGT	CCGAGAGAGG	TCACCTTACTT	GCTGCTGCTT
1090	1100	1110	1120	1130
AAAACGAGGA	AGATGCACAC	CATGTACAAG	AAACCAGACA	CCCAGACAAT
1150	1160	1170	1180	1190
COITCAGAGT	TTAACTCGTT	CGTCATCCCG	AGCCTATGGT	CTACAGGCCT
1210	1220	1230	1240	1250
GTEAGATCAC	GCATTAAGAT	GCTTTTGGCC	AAGAAGACCA	AGCGAGAGTT
1270	1280	1290	1300	1310
CTCGACGCGT	CGTCAGCCAG	GGATGCTGAA	CAAGAGGAGA	AGGAGAGGTT
1330	1340	1350	1360	1370
CTGACTAGAG	AAGCCTTACC	ACCCCTCGTC	CCCATCGCGC	CGCGGAGAC
1390	1400	1410	1420	1430
GACGTCGACG	TTGAAGAACT	AGAGTATCAC	GCAGGTGCAG	GGGTCGTGGA
1450	1460	1470	1480	1490
AGCGCGTTGA	AAGTCACCGC	ACAGCCGAAC	GACGTACTAC	TAGGAAATTA
1510	1520	1530	1540	1550
TCCCGCAGCA	CCGTGCTCAA	GAGCTCCAAG	TTGGCCCCCG	TGCACCCTCT
1570	1580	1590	1600	1610
GTGAAAAATA	TAACACATAA	CGGGAGGGCC	GGCGGTACC	AGGTCGACGG
1630	1640	1650	1660	1670
AGGGTCCTAC	TACCATGTGG	ATCGGCCATT	CCGGTCCCTG	AGTTTCAAGC
1690	1700	1710	1720	1730
AGCGCCACTA	TGGTGATCAA	CGAAAGGGAG	TTCGTCAACA	GGAAACTATA
1750	1760	1770	1780	1790
				1800

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GTTCACGGAC	CGTCGCTGAA	CAGTACGAG	GAGAACTACG	AGAAAGTCAG	AGCTTGA
1810	1820	1830	1840	1850	1860
ACTGACGCCG	AGTACGTGTT	CGACGTAGAT	AAAAAATGCT	GCGTCAAGAG	AGAGGAAGCG
1870	1880	1890	1900	1910	1920
TCGGGTITGG	TGTTGGTGGG	AGAGCTAACC	AACCCCCCGT	TCCATGAATT	CGCCTACGAA
1930	1940	1950	1960	1970	1980
GGGCTGAAGA	TCAGGCCGTC	GGCACCATAT	AAGACTACAG	TAGTAGGAGT	CTTTGGGGTT
1990	2000	2010	2020	2030	2040
CCGGGATCAG	GCAAGTCTGC	TATTATTAAG	AGCCTCGTGA	CCAAACACGA	TCTGGTCACC
2050	2060	2070	2080	2090	2100
AGCGGCAAGA	AGGAGAACTG	CCAGGAAATA	GTTAACGACG	TGAAGAAGCA	CCGCGGGAAG
2110	2120	2130	2140	2150	2160
GGGACAAGTA	GGGAAAACAG	TGACTCCATC	CTGCTAAACG	GGTGTGCTCG	TGCCGTGGAC
2170	2180	2190	2200	2210	2220
ATCCTATATG	TGGACGAGGC	TTTCGCTTGC	CATTCCGGTA	CTCTGCTGGC	CCTAATTGCT
2230	2240	2250	2260	2270	2280
CTTGTTAAAC	CTCGGAGCAA	AGTGGTGTTA	TGCGGAGACC	CCAAGCAATG	CGGATTCTTC
2290	2300	2310	2320	2330	2340
AATATGATGC	AGCTTAAGGT	GAACITCAAC	CACAACATCT	GCACTGAAGT	ATGTCATAAA
2350	2360	2370	2380	2390	2400
AGTATATCCA	GACGTTGCAC	GCGTCCAGTC	ACGGCCATCG	TGTCTACGTT	GCACTACGGA
2410	2420	2430	2440	2450	2460
GGCAAGATGC	GCACGACCAA	CCCGTGCAAC	AAACCCATAA	TCATAGACAC	CACAGGACAG
2470	2480	2490	2500	2510	2520
ACCAAGCCCA	AGCCAGGAGA	CATCGTGTTA	ACATGCTTCC	GAGGCTGGGC	AAAGCAGCTG
2530	2540	2550	2560	2570	2580
CAGTTGGACT	ACCGTGGACA	CGAAGTCATG	ACAGCAGCAG	CATCTCAGGG	CCTCACCCGC
2590	2600	2610	2620	2630	2640
AAGGGGTAT	ACGCCGTAAG	GCAGAAGGTG	AATGAAATC	CCTTGATATG	CCCTGCGTCG
2650	2660	2670	2680	2690	2700
GAGCACGTGA	ATGTACTGCT	GACGCGCACT	GAGGATAGGC	TGGTGTGGAA	AACGCTGGCC
2710	2720	2730	2740	2750	2760
GGGATCCCT	GGATTAAGGT	CCTATCAAAC	ATTCACACGG	GTAACCTTAC	GGCCACATTG
2770	2780	2790	2800	2810	2820
GAAGAATGGC	AAGAAGAACA	CGACAAAATA	ATGAAGGTGA	TTGAAGGACC	GGCTGCGCCT
2830	2840	2850	2860	2870	2880
GTCGACGCGT	TCCAGAACAA	AGCGAACGTG	TGTTGGGCGA	AAAGCCTGGT	GCCTGTCCTG
2890	2900	2910	2920	2930	2940
GAACTGCCG	GAATCAGATT	GACAGCAGAG	GAGTGGAGCA	CCATAATTAC	AGCATTTAAG
2950	2960	2970	2980	2990	3000
GAGGACAGAG	CTTACTCTCC	AGTGGTGGCC	TTGAATGAAA	TTTGACCCAA	GTACTATGGA
3010	3020	3030	3040	3050	3060
GTTGACCTGG	ACAGTGGCCT	GTTTTCTGCC	CCGAAGGTGT	CCCTGTATTIA	CGAGAACAAC
3070	3080	3090	3100	3110	3120
CACTGGGATA	ACAGACCTGG	TGGAAGGATG	TATGGATTCA	ATGCCGCAAC	AGCTGCCAGG
3130	3140	3150	3160	3170	3180
CTGGAAGCTA	GACATACCTT	CCTGAAGGGG	CAGTGGCATA	CGGGCAAGCA	GGCAGTTATC
3190	3200	3210	3220	3230	3240
GCAGAAAGAA	AAATCCAACC	GCTTTCGTG	CTGGACAATG	TAATTCCTAT	CAACCGCAGG
3250	3260	3270	3280	3290	3300
CTGCCGACG	CCCTGGTGGC	TGAGTACAAG	ACGGTTAAAG	GCAGTAGGGT	TGAGTGGCTG
3310	3320	3330	3340	3350	3360
GTCAATAAAG	TAAGAGGGTA	CCACGTCCTG	CTGGTGAGTG	AGTACAACCT	GGCTTTGCCT
3370	3380	3390	3400	3410	3420
CGACGCAGGG	TCACTTGTTT	GTCACCGCTG	AATGTCACAG	GCGCCGATAG	GTGCTACGAC
3430	3440	3450	3460	3470	3480
CTAAGTTTAG	GACTGCCGGC	TGACGCCGGC	AGGTTTCGACT	TGGTCTTTGT	GAACATTAC
3490	3500	3510	3520	3530	3540
ACGGAATTCA	GAATCCACCA	CTACCAGCAG	TGTGTCGACC	ACGCCATGAA	GCTGCAGATG

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3550	3560	3570	3580	3590	3600
CTTGGGGGAG	ATGCGCTACG	ACTGCGCTAA	CCCGGCGGCA	TCTTGATGAG	AGCTTACG
3610	3620	3630	3640	3650	3660
TACGCCGATA	AAATCAGCGA	AGCCGTTGTT	TCCTCCCTAA	GCAGAAAGTT	CTCGTCTGCA
3670	3680	3690	3700	3710	3720
AGAGTGTTCG	GCCCGGATTG	TGTCACCAGC	AATACAGAAG	TGTTCTTGCT	GTTCTCCAAC
3730	3740	3750	3760	3770	3780
TTTGACAACG	GAAAGAGACC	CTCTACGCTA	CACCAGATGA	ATACCAAGCT	GAGTGCCGTG
3790	3800	3810	3820	3830	3840
TATGCCGGAG	AAGCCATGCA	CACGGCCGGG	TGTGCACCAT	CCTACAGAGT	TAAGAGAGCA
3850	3860	3870	3880	3890	3900
GACATAGCCA	CGTGACACAG	AGCGGCTGTG	GTAAACGCAG	CTAACGCCCG	TGGAAGTGTG
3910	3920	3930	3940	3950	3960
GGGGATGGCG	TATGCAGGGC	CGTGGCGAAG	AAATGGCCGT	CAGCCTTTAA	GGGAGCAGCA
3970	3980	3990	4000	4010	4020
ACACCAGTGG	GCACAATTAA	AACAGTCATG	TGCGGCTCGT	ACCCCGTCAT	CCACGCTGTA
4030	4040	4050	4060	4070	4080
GCGCCTAATT	TCTCTGCCAC	GACTGAAGCG	GAAGGGGACC	GCGAATTGGC	CGCTGTCTAC
4090	4100	4110	4120	4130	4140
CGGGCAGTGG	CCGCCGAAGT	AAACAGACTG	TCCTGAGCA	GCGTAGCCAT	CCCGCTGCTG
4150	4160	4170	4180	4190	4200
TCCACAGGAG	TGTTACGCGG	CGGAAGAGAT	AGGCTGCAGC	AATCCCTCAA	CCATCTATTG
4210	4220	4230	4240	4250	4260
ACAGCAATGG	ACGCCACGGA	CGCTGACGTG	ACCATCTACT	GCAGAGACAA	AAGTTGGGAG
4270	4280	4290	4300	4310	4320
AAGAAAATCC	AGGAAGCCAT	TGACATGAGG	ACGGCTGTGG	AGTTGCTCAA	TGATGACGTG
4330	4340	4350	4360	4370	4380
GAGCTGACCA	CAGACTTGGT	GAGAGTGCAC	CCGGACAGCA	GCCTGGTGGG	TCGTAAGGGC
4390	4400	4410	4420	4430	4440
TACAGTACCA	CTGACGGGTC	GCTGTACTCG	TACTTTGAAG	GTACGAAATT	CAACCAAGCT
4450	4460	4470	4480	4490	4500
GCTATTGATA	TGGCAGAGAT	ACTGACGTTG	TGGCCCAGAC	TGCAAGAGGC	AAACGAACAG
4510	4520	4530	4540	4550	4560
ATATGCGCTAT	ACGCGCTGGG	CGAAACAATG	GACAACATCA	GATCCAAATG	TCCGGTGAAC
4570	4580	4590	4600	4610	4620
GATTCGATT	CATCAACACC	TCCCAGGACA	GTGCCCTGCC	TGTGCCGCTA	CGCAATGACA
4630	4640	4650	4660	4670	4680
GCAGAACGGA	TGCCCCGCTT	TAGGTACAC	CAAGTTAAAA	GCATGGTGGT	TTGCTCATCT
4690	4700	4710	4720	4730	4740
TTTCCCTCC	CGAAATACCA	TGTAGATGGG	GTGCAGAAGG	TAAAGTGCGA	GAAGGTTCTC
4750	4760	4770	4780	4790	4800
CTGTTGACG	CGACGGTACC	TTCAGTGGTT	AGTCCGCGGA	AGTATGCCGC	ATCTACGACG
4810	4820	4830	4840	4850	4860
GACCACTCAG	ATCGGTGCTT	ACGAGGGTTT	GACTTGGACT	GGACCACCGA	CTCGTCTTCC
4870	4880	4890	4900	4910	4920
ACTGCCAGCG	ATACCATGTC	GCTACCCAGT	TTGCAGTCGT	GTGACATCGA	CTCGATCTAC
4930	4940	4950	4960	4970	4980
GAGCCAATGG	CTCCCATAGT	AGTGACGGCT	GACGTACACC	CTGAACCCGC	AGGCATCGCG
4990	5000	5010	5020	5030	5040
GACCTGGCGG	CAGATGTGCA	CCCTGAACCC	GCAGACCATG	TGGACCTCGA	GAACCCGATT
5050	5060	5070	5080	5090	5100
CCTCCACCGC	GCCCGAAGAG	AGCTGCATAC	CTTGCCCTCC	GCGCGGCGGA	GCGACCGGTG
5110	5120	5130	5140	5150	5160
CCGGCGCCGA	GAAAGCCGAC	GCCTGCCCCA	AGGACTGCGT	TTAGCAACAA	GCTGCCTTTG
5170	5180	5190	5200	5210	5220
ACGTTGCGCG	ACTTTGACGA	GCACGAGGTC	GATGCGTTGG	CCTCCGGGAT	TACTTTCCGA
5230	5240	5250	5260	5270	5280
GACTTCGACG	ACGTCCTGCC	ACTAGCCCGC	GCGGGTGCAT	ATATTTTCTC	CTCGGACACT
5290	5300	5310	5320	5330	5340

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GGCAGCGGAC	ATTTACAACA	AAATCGTT	AGGCAGCACA	ATCTCCAGTG	CGCAATG
5350	5360	5370	5380	5390	5400
GATGCGGTCC	AGGAGGAGAA	AATGTACCCG	CCAAAATTGG	ATACTGAGAG	GGAGAAGCTG
5410	5420	5430	5440	5450	5460
TTGCTGCTGA	AAATGCAGAT	GCACCCATCG	GAGGCTAATA	AGAGTCGATA	CCAGTCTCGC
5470	5480	5490	5500	5510	5520
AAAGTGAGGA	ACATGAAAGC	CAOGGTGGTG	GACAGGCTCA	CATCGGGGGC	CAGATTGTAC
5530	5540	5550	5560	5570	5580
ACGGGAGCGG	ACGTAGGCCG	CATACCAACA	TACGCGGTTC	GGTACCCCCG	CCCCGTGTAC
5590	5600	5610	5620	5630	5640
TCCCCTACCG	TGATCGAAAG	ATTCTCAAGC	CCCGATGTAG	CAATCGCAGC	GTGCAACGAA
5650	5660	5670	5680	5690	5700
TACCTATCCA	GAAATTACCC	AACAGTGGCG	TCGTACCAGA	TAACAGATGA	ATACGACGCA
5710	5720	5730	5740	5750	5760
TACTTGAGCA	TGGTTGACGG	GTCGGATAGT	TGCTTGAGCA	GAGCGACATT	CTGCCCGGCG
5770	5780	5790	5800	5810	5820
AAGCTCCGGT	GCTACCCGAA	ACATCATGCG	TACCACCAGC	CGACTGTACG	CAGTGCCGTC
5830	5840	5850	5860	5870	5880
CCGTACCCCT	TTCAGAACAC	ACTACAGAAC	GTGCTAGCGG	CGCCACCAA	GAGAAACATC
5890	5900	5910	5920	5930	5940
AACGTCACGC	AAATGCGAGA	ACTACCCACC	ATGGACTCGG	CAGTGTTCAG	CGTGGAGTGC
5950	5960	5970	5980	5990	6000
TTCAAGCGCT	ATGCCTGCTC	CGGAGAATAT	TGGGAAGAAT	ATGCTAAACA	ACCTATCOGG
6010	6020	6030	6040	6050	6060
ATAACCACTG	AGAACATCAC	TACCTATGTG	ACCAAAATTGA	AAGGCCCGAA	AGCTGCTGCC
6070	6080	6090	6100	6110	6120
TTGTTGCTA	AGACCCACAA	CTTGGTTCCG	CTGCAGGAGG	TTCCCATGGA	CAGATTCAAG
6130	6140	6150	6160	6170	6180
GTCGACATGA	AACGAGATGT	CAAGTCACT	CCAGGGACGA	AACACACAGA	GGAAAGACCC
6190	6200	6210	6220	6230	6240
AAAGTCCAGG	TAATTCAAGC	AGCGGAGCCA	TTGGCGACCG	CTTACCTGTG	CGGCATCCAC
6250	6260	6270	6280	6290	6300
AGCGAATTAG	TAAGGAGACT	AAATGCTGTG	TTACGCCCTA	ACGTGCACAC	ATTGTTTGAT
6310	6320	6330	6340	6350	6360
ATGTCGGCCG	AAGACTTTGA	CGCGATCATC	GCCTCTCACT	TCCACCCAGG	AGACCCGGTT
6370	6380	6390	6400	6410	6420
CTAGAGACGG	ACATTGCATC	ATTTCGACAA	AGCCAGGACG	ACTCCTTGCC	TCTTACAGGT
6430	6440	6450	6460	6470	6480
TTAATGATCC	TCAAGATCT	AGGGGTGGAT	CAGTACCTGC	TGGACTTGAT	CGAGGCAGCC
6490	6500	6510	6520	6530	6540
TTTGGGGAAA	TATCCAGCTG	TCACCTACCA	ACTGGCACGC	GCTTCAAGTT	CGGAGCTATG
6550	6560	6570	6580	6590	6600
ATGAAATCGG	GCATGTTTCT	GACTTTGTTT	ATTAAACTG	TTTTGAACAT	CACCATAGCA
6610	6620	6630	6640	6650	6660
AGCAGGGTAC	TGGAGCAGAG	ACTCACTGAC	TCCGCCTGTG	CGGCCTTCAT	CGGCGACGAC
6670	6680	6690	6700	6710	6720
AACATCGTTC	ACGGAGTGAT	CTCCGACAAG	CTGATGGCGG	AGAGGTGCGC	GTCGTGGGTC
6730	6740	6750	6760	6770	6780
AACATGGAGG	TGAAGATCAT	TGACGCTGTC	ATGGGCGAAA	AACCCCCATA	TTTTGTGGG
6790	6800	6810	6820	6830	6840
GGATTATAG	TTTTTGACAG	CGTCACACAG	ACCGCCTGCC	GTGTTTCAGA	CCCACTTAAG
6850	6860	6870	6880	6890	6900
CGCCTGTTCA	AGTTGGGTAA	GCCGCTAACA	GCTGAAGACA	AGCAGGACGA	AGACAGGCGA
6910	6920	6930	6940	6950	6960
CGAGCACTGA	GTGACGAGGT	TAGCAAGTGG	TTCCGGACAG	GCTTGGGGGC	CGAACTGGAG
6970	6980	6990	7000	7010	7020
GTGGCACTAA	CATCTAGGTA	TGAGGTAGAG	GGCTGCAAAA	GTATCCTCAT	AGCCATGGCC
7030	7040	7050	7060	7070	7080
ACCTTGCGCA	GGGACATTAA	GGCGTTTAAG	AAATTGAGAG	GACCTGTTAT	ACACCTCTAC

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7090	7100	7120	7130	7140
GGCGGTCCTA	GATTGGTGGC	TTAATACCAA	GAATTCTGAT	TGGATCCCGG
GTAATTAA				
7150	7160	7170	7180	7190
7200				
GAATTACATC	CCTACGCAAA	CGTTTTACGG	CCGCCCGGTGG	CGCCCGCGCC
CGGCGGCCCCG				
7210	7220	7230	7240	7250
7260				
TCCTTGGCCG	TTGCAGGCCA	CTCCGGTGGC	TCCCGTCTGC	CCCGACTTCC
AGGCCAGCA				
7270	7280	7290	7300	7310
7320				
GATGCAGCAA	CTCATCAGCG	CCGTAAATGC	GCTGACAATG	AGACAGAACG
CAATTGCTCC				
7330	7340	7350	7360	7370
7380				
TGCTAGGCCT	CCCAAACCAA	AGAAGAAGAA	GACAACCAAA	CCAAAGCCGA
AAACGCAGCC				
7390	7400	7410	7420	7430
7440				
CAAGAAGATC	AACGGAAAAA	CGCAGCAGCA	AAAGAAGAAA	GACAAGCAAG
CCGACAAGAA				
7450	7460	7470	7480	7490
7500				
GAAGAAGAAA	CCCGGAAAAA	GAGAAAGAAT	GTGCATGAAG	ATTGAAAATG
ACTGTATCTT				
7510	7520	7530	7540	7550
7560				
CGTATGCGGC	TAGCCACAGT	AACGTAGTGT	TTCCAGACAT	GTCGGGCACC
GCACTATCAT				
7570	7580	7590	7600	7610
7620				
GGGTGCAGAA	AATCTCGGGT	GGTCTGGGGG	CCTTCGCAAT	CGGCGCTATC
CTGGTGCTGG				
7630	7640	7650	7660	7670
7680				
TTGTGGTCAC	TTGCATTGGG	CTCCGCAGAT	AAGTTAGGGT	AGGCAATGGC
ATTCATATAG				
7690	7700	7710	7720	7730
7740				
CAAGAAAATT	GAAAACAGAA	AAAGTTAGGG	TAAGCAATGG	CATATAACCA
TAACTGTATA				
7750	7760	7770	7780	7790
7800				
ACTTGTAACA	AAGCGCAACA	AGACCTGCGC	AATTGGCCCC	GTGGTCCGCC
TCACGGAAAC				
7810	7820	7830	7840	7850
7860				
TCGGGGCAAC	TCATATTGAC	ACATTAATTG	GCAATAATTG	GAAGCTTACA
TAAGCTTAAT				
7870	7880	7890	7900	7910
7920				
TGGACGAATA	ATTGGATTAT	TATTTTATTT	TGCAATTGGT	TTTTAATATT
TCCAAAAAAA				
7930	7940	7950	7960	7970
7980				
AAAAAAAAAA	AAAAAAAAAA	AAAAAAAAAA	AAAAAAAAAA	AAAAAAAAAA
AAAAAAAAAA				
7990	8000	8010	8020	8030
8040				
AAA.....

